

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
21 February 2002 (21.02.2002)

PCT

(10) International Publication Number
WO 02/13702 A1

(51) International Patent Classification⁷: **A61B 17/064** (74) Agent: **HIENZ, William, M., III**; Dicke, Billig & Czaja, P.A., Suite 1250, 701 Fourth Avenue South, Minneapolis, MN 55415 (US).

(21) International Application Number: **PCT/US01/07337**

(22) International Filing Date: 7 March 2001 (07.03.2001) (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/187,428 7 March 2000 (07.03.2000) US
60/201,594 3 May 2000 (03.05.2000) US

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(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

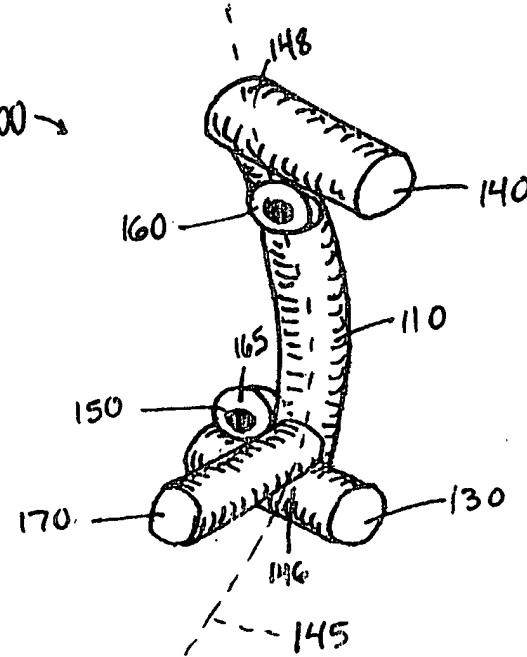
Published:
— with international search report

[Continued on next page]

(54) Title: EVERTING STAPLE DEVICES AND METHODS



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(57) Abstract: A surgical staple for use in creating an everted anastomosis of at least two anatomical structures includes a bendable staple body. At least two evertting elements protrude from the staple body. Spacing elements and a penetrating element are also provided. In use, the evertting elements and spacing elements are apposed, and inner layers of the anatomical structures are held together to form an everted anastomosis. The evertting platforms are disposed on an outer radius of the closed staple, and the penetrating element is disposed on a different, inner radius, to ensure that the penetrating element is excluded from the lumen of the anastomosis. Embodiments of the invention substantially ensure intima-to-intima approximation completely circumferentially at the anastomotic site, with no portion of cut tissue edges, suture or staple exposed to the lumen of the anastomosis. Cut tissue edges, as well as the staple, are completely extra-luminal. Related methods are also disclosed.

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EVERTING STAPLE DEVICES AND METHODS

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CROSS-REFERENCE TO RELATED APPLICATIONS

The subject matter of this application is related to the subject matter of commonly assigned U.S. Applications Nos. 60/187,428, filed March 7, 2000 and 10 60/201,594, filed May 3, 2000, both of which are incorporated herein by reference and priority to which is claimed under 35 U.S.C. § 119(e).

BACKGROUND OF THE INVENTION

1. Field of Invention

15 The invention relates to devices and methods for anastomosing two or more anatomical structures, such as vascular structures. Embodiments according to the invention are desirable in both minimally invasive and conventional surgical situations. Certain embodiments have particular application to minimally invasive direct coronary artery bypass (MIDCAB) and off-pump 20 (OPCAB) procedures, for example.

2. Description of Related Art

Surgical connections (anastomoses) between two tubular structures of soft tissue (e.g. blood vessels) typically are fashioned with "linkage" devices, 25 such as flexible suture or rigid staples. Such anastomoses involve surgical incisions of the two soft tissue structures, which result in cut edges. These edges ideally are excluded from the inside channel (lumen) of the two joined structures. Ideally, the linkage devices are also excluded from the lumen.

The suture- and staple-type linkage devices of the prior art, however, do not always achieve the above-described ideal; cut tissue edges and the linkage devices themselves sometimes are exposed to the lumen. See Figures 1A-1D, for example, which show unpredictable eversion and exclusion of cut edges 10 5 (of ends 20, 30 of anatomical structures 40, 50) and suture 60 from lumen 70 when sutures alone are used, as in the prior art.

SUMMARY OF THE INVENTION

In view of these and other disadvantages, a need exists for devices and 10 methods that consistently, predictably and reliably evert the edges of an e.g. circumferential anastomosis between two tubular anatomical structures, e.g. a vascular anastomosis between the left internal mammary artery and the left anterior descending coronary artery. Such devices and methods can be used alone, or in conjunction with a biological adhesive product, suture, or other 15 supplemental products or devices. They can be part of a sutured or stapled anastomosis, i.e. an anastomosis using one or more sutures and/or staples in combination with the devices of the invention.

The ultimate purpose, according to embodiments of the invention, is to substantially ensure inner-layer-to-inner-layer (e.g. intima-to-intima) 20 approximation completely circumferentially at the anastomotic site, with no portion of the cut edges of the tissues or suture or staple exposed to the lumen of the anastomosis. Instead, embodiments of the invention substantially ensure that the cut edges of the tissues, as well as the linkage device, are completely extra-luminal. With such embodiments, the entire circumference of both structures of 25 the anastomosis is completely everted.

Embodiments of the invention provide a number of advantages. For example, the invention provides a more precise, completely everted anastomosis more predictably than heretofore possible with staples or suture alone.

Embodiments of the invention are believed to result in a higher patency rate in 5 vascular anastomoses, by eliminating cut edges of tissue or foreign bodies (e.g. sutures or staples) from the lumen. Such tissue edges or foreign bodies, if left exposed to the lumen, can threaten the patency of the anastomosis by allowing thrombus (clot) formation and/or fibrosis or intimal hyperplasia (scarring), ultimately resulting in stenosis (narrowing) or occlusion. Embodiments of the 10 invention, on the other hand, cause immediate and long-term improvements in patency and therefore quality of life and longevity.

Further, embodiments of the invention allow a smaller number of anastomotic eversion devices to be used, e.g. in conjunction with a biological adhesive or equivalent. Such hybrid procedures allow for a simpler, more 15 expeditious anastomosis, with either conventional or minimally invasive techniques, with or without the aid of cardiopulmonary bypass.

Still further, according to embodiments of the invention, everting platforms on an outer radius of a closed evertng staple assure that a penetrating element of the staple is excluded from the lumen, because the penetrating 20 element is on a different, inner radius. Spacing elements according to the invention assure a space between everting platforms of the staple in a closed configuration of the staple, negating any tissue necrosis at the level of intima-to-intima approximation.

Penetrating and spacing elements according to embodiments of the 25 invention may cause tissue damage, but such damage is outside of the

anastomosis, i.e. it is at or near the inner radius of the closed staple.

Nevertheless, spacing elements and/or penetrating elements according to such embodiments have small surface area, to minimize tissue necrosis. To effect parallel alignment of the evertting platforms, the staple is "scissored" in the 5 closed position thereof. One of the evertting platforms is offset accordingly.

By providing two evertting platforms on a single piece of wire or other material constituting the staple, according to embodiments of the invention, the need to deliver an opposing evertting platform from an opposite side of an anastomosis is eliminated. Other features and advantages according to 10 embodiments of the invention will become apparent from the remainder of this application.

BRIEF DESCRIPTION OF THE DRAWINGS

Embodiments of the invention will be described with reference to the 15 figures, in which like reference numerals refer to like elements and in which:

Figures 1A - 1D show one type of prior art anastomosis technique using a suture alone, with unpredictable eversion and exclusion of cut edges and suture from the lumen;

Figure 2 is a panoramic view of an open staple according to an 20 embodiment of the invention, with an evertting platform, offset evertting platform, two spacing elements and penetrating element as shown;

Figure 3 is a panoramic view of a closed, "scissored" staple with two radii and two axes according to an embodiment of the invention. More specifically, Figure 3 shows an outer radius of the illustrated evertting platforms,

an inner radius of the spacing and penetrating elements, and side-by-side offset axes of the closed staple;

Figure 4 is a side view of a staple according to an embodiment of the invention, showing the penetrating element, spacing elements, and everting platforms;

Figure 5 is a side view of closed staple enclosing tissue, according to an embodiment of the invention, showing the penetrating element, everting platforms and spacing elements;

Figure 6 is a side view showing the inner radius of the spacing and 10 penetrating elements and the outer radius of the everting platforms;

Figure 7 is an end view of a closed staple enclosing tissue, according to an embodiment of the invention, showing the penetrating element, everting platforms and spacing elements;

Figure 8 is an inside view of a open staple, according to an embodiment 15 of the invention, showing the penetrating element, everting platforms and spacing elements, and the direction of closure of the staple;

Figure 9 is an outside view of an open staple, according to an embodiment of the invention, showing the penetrating element, everting platforms and spacing elements, and the direction of closure of the staple;

20 Figure 10 is a perspective view of a staple and an eversion platform, according to an embodiment of the invention;

Figure 11 is a side view of an eversion platform with overhanging collar, according to an embodiment of the invention;

25 Figure 12 is a side, cross-sectional view of a stapled anastomosis in a staple-open, platforms-apart configuration;

Figure 13 is a side, cross-sectional view of a stapled anastomosis in a staple-closed, platforms-apposed configuration;

Figure 14 is a side view showing the inner radius of the staple and the outer radius of the eversion platforms; and

5 Figure 15 is a panoramic view of an end-to-side anastomosis in a staple-closed configuration.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

10 A first embodiment of surgical staple 100 for creating a completely everted anastomosis is shown in Figures 2-9. Staple body 110 of surgical staple 100 is constructed to bend, e.g. at 120 in Figure 3. At opposite ends of staple body 110, staple 100 includes evertting platforms or elements 130, 140 that protrude from staple body 110. Evertting platform 130 is centered with respect to 15 longitudinal axis 145 of staple body 110. As illustrated, a central portion 146 of evertting platform 130 is connected to the very end of staple body 110. Evertting platform 140, on the other hand, is connected to staple body 110 at its end 148, not at a central portion thereof. Thus, as shown in e.g. Figures 8-9, evertting platforms 130, 140 are offset with respect to each other, in a transverse direction 20 with respect to axis 145.

Surgical staple 100 also includes first spacing element 150, disposed adjacent staple body 110 at the very end thereof. Specifically, spacing element 150 is adjacent to longitudinal axis 145 thereof and above evertting platform 130, as viewed in e.g. Figures 2 and 8. Staple 100 also includes second spacing 25 element 160, disposed along staple body 110 at the opposite end thereof and generally in line with longitudinal axis 145 thereof. Spacing elements 150, 160

are closer to the middle of staple body 100 than are everting platforms 130, 140.

Spacing elements 150, 160 can be generally tapered, as shown at e.g. 165 in Figures 2 and 8. Of course, other placements and shapes of spacing elements 150, 160 are contemplated.

5 Penetrating element 170 is constructed to penetrate the anatomical structures being anastomosed, in a manner to be described, and is disposed at one end of staple body 110. Penetrating element 170 is disposed generally along longitudinal axis 145 of staple body 110 and is generally perpendicular or transverse, or otherwise angled, to it.

10 Staple 100 is constructed to bend from the straight or non-use or open configuration illustrated in Figures 2 and/or 4 to the bent or "scissored" or closed configuration of Figure 3. In the Figure 3 configuration, everting platforms 130, 140 are disposed generally parallel to each other on the same side of penetrating element 170, e.g. the underside as viewed in Figure 3. Additionally, everting platforms 130, 140 are apposed, as are spacing elements 150, 160.

15 As illustrated in Figure 3, spacing elements 150, 160 and penetrating element 170 generally define elevation line 180. Evertng platforms 130, 140 generally define elevation line 190. Lines 180, 190 define distances from bend 120 that can be considered inner and outer radii, respectively. Because line 180 20 is closer to bend 120, line 180 can be considered to define an inner radius of staple 100. Similarly, line 190 can be considered to define an outer radius of staple 100. Additionally, Figure 3 illustrates side-by-side offset axes 200, 210 of staple 100. Axes 200, 210 are offset from each other in two dimensions, e.g. a first dimension as viewed in the direction of lines 180, 190, and a second 25 dimension as viewed in the direction of evertng platforms 130, 140, in Figure 3.

In its closed configuration, staple 100 is bent in a "U" shape to form two legs 220, 230 separated by bend 120. Legs 220, 230 are disposed along axes 200, 210 and thus are offset from each other in two dimensions.

Figures 5-7 show staple 100 in use. As shown, staple 100 joins inner 5 layer (intima) 240 of anatomical structure 250 to inner layer (intima) 260 of anatomical structure 270. Staple 100 keeps lumen 280 between anatomical structures 250, 270 free of exposure to cut tissue edges 290, ends 300, 310 of anatomical structures 250, 270, and all portions of staple 100 itself, including 10 penetrating element 170. Lumen 280 is also free of suture and other foreign bodies. Thus, staple 100 causes immediate and long-term improvements in the 15 patency of the anastomosis by reducing the likelihood of clot formation, scarring, stenosis and other complications.

When staple 100 is in its closed configuration around cut ends 300, 310 of anatomical structures 250, 270, spacing elements 150, 160 are in an apposed 15 relationship and cut ends 300, 310 are held together between spacing elements 150, 160. Additionally, everting platforms 130, 140 are in an apposed relationship, with cut ends 300, 310 held together between everting platforms 130, 140. Penetrating element 170 penetrates ends 300, 310. The distance 20 between apposed spacing elements 150, 160 is less than the distance between apposed everting platforms 130, 140, as shown. As shown in e.g. Figure 6, apposed spacing elements 150, 160 and/or penetrating element 170 define inner radius 180 of closed staple 100, and apposed everting platforms 130, 140 define outer radius 190 of closed staple 100, radii 180, 190 being considered to originate at bend 120 in staple 100. Because penetrating element 170 is on the

different, inner radius 180, it is generally assured that penetrating element 170 is excluded from lumen 280.

As shown in Figures 8-9, evertng platform 140 moves laterally or transversely with respect to staple body 110 as staple 100 is closed, such that the 5 direction of closure 320 is at an angle to staple body 110. A method of bending a surgical staple according to an embodiment of the invention includes moving evertng platform 140, disposed at one end of staple 100, to be parallel to and adjacent to evertng platform 130 disposed at an opposite end of staple 100.

Spacing element 160 disposed at one end of staple 100 becomes apposed to 10 spacing element 150 disposed at the opposite end of staple 100. Spacing elements 150, 160 are disposed adjacent to and generally parallel to penetrating element 170. Evertng platform 140 moves in direction of closure 320 to its disposition parallel to and adjacent to evertng platform 130. Direction of closure 320 is disposed at an angle, e.g. about 15° to about 20°, to longitudinal 15 axis 145 extending between evertng platforms 130, 140. Thus, evertng platforms 130, 140 are moved from an offset configuration, illustrated in Figure 8, in which platforms 130, 140 are offset from each other along their respective longitudinal axes, to a non-offset configuration, illustrated in e.g. Figure 3, in which evertng platforms 130, 140 are aligned with each other.

20 Turning to Figures 10-15, staple devices according to embodiments of the invention include two main parts. The first part is staple 400, defining a generally curved shape and supporting a fixed eversion platform 410 securely and immovably affixed thereto, according to one embodiment. Staple 400 can be a free-standing staple, or attached to a suture or other delivery system, for

example. The second part is free eversion platform 420, which is supportable by staple 400 and slidably relative thereto in a "threaded" relationship.

Both staple 400 and free eversion platform 420 preferably have a substantially "D" - shaped cross section 430, to predictably orient eversion platform 420. Of course, other cross-sectional shapes of the staple and free eversion platform are contemplated, according to embodiments of the invention. 5
Staple 400 and platform 420 preferably are made of a biocompatible material. At least the staple is malleable, according to a preferred embodiment, so that it can be crimped into a closed position with a closing or crimping device. 10
According to other embodiments, described below, a memory metal eliminates or reduces the need for a separate closing device.

Free eversion platform 430 generally comprises two parts: collar or spacing element 440 and eversion body or platform 450. Eversion platform 450 preferably is made of a biocompatible material similar to staple 400. However, 15 it is substantially non-deformable, according to this embodiment. Collar 440 slightly overhangs eversion platform 420 in the axis of staple 460, as indicated at 460 in Figure 11. Overhang 460 substantially prevents apposing eversion platforms 410, 420 from crushing cut ends 470, 480 of vascular or other structures 490, 500 in the anastomosis, by keeping them slightly separated from 20 one another in the closed position. Thus, any tissue in the anastomosis is approximated without being strangulated. Additionally, lumen 510 is kept free of exposure to cut tissue edges 520, 530, staple 400, eversion platforms 410, 420, suture, or other structures that might threaten the patency of the anastomosis.

According to embodiments of the invention, either eversion platform 410, 420 illustrated in e.g. Figure 10 can be fixed or free, or both can be free. Everting staple 400 likely is easier to deploy, however, if one platform is fixed and the other is free, as illustrated. Structurally, platforms 410, 420 are generally identical according to one embodiment.

According to the illustrated embodiment, fixed eversion platform 410 is nearer trailing end 535 of staple 400, especially when it is to be deployed via a suture or other delivery mechanism. Suture can be attached to the leading end 537 of the staple, for example, and free eversion platform 420 deployed after 10 engaging and penetrating the two vascular or other structures 490, 500 with staple 400. The body of staple 400, or a portion thereof, thus is a penetrating element. According to one embodiment, free eversion platform 420 is threaded over a needle and suture, is advanced along the suture to staple 400, and then is advanced to a desired position along staple 400 as shown in e.g. Figure 12. A 15 closing/crimping device or other means then brings the paired eversion platforms 410, 420 into final juxtaposition to each other as staple 400 is closed, in the manner of e.g. Figure 13. Eversion platforms 410, 420 are the fulcrum for closing staple 400.

Of course, two separate free eversion platforms 410 can be employed, 20 with neither affixed to staple 400 at least until after staple 400 engages the two vascular structures 490, 500. The free eversion platforms slide over suture and/or the respective ends of staple 400, while staple 400 is still in an open position. According to this embodiment, and/or according to those described earlier, suture can be attached to either or both ends of staple 400.

As shown in Figure 14, collars 440 and eversion bodies 450 of eversion platforms 410, 420 define inner radius 540 and outer radius 550, respectively, of staple 400. Figure 15 shows multiple closed staples 400 disposed in a radial plane, with eversion bodies 450 disposed in the anastomotic plane, in an end-to-side anastomosis.

According to embodiments of the invention, a plurality of evertting staples can be employed to create a single anastomosis, as shown in Figure 15, by themselves or optionally in combination with supplemental adhesives, suture, etc. to reduce the number of staples needed. Of course, each eversion platform and staple can be appropriately dimensioned to suit a particular patient, surgical procedure, surgical environment or other factor. Device pairs or combinations of 10 different types or sizes can be used in the same anastomosis, if desired.

As referenced earlier, staple devices according to the multiple embodiments of the invention can be free-standing staples and/or can be 15 connected to a delivery system for the staple device. For example, a flexible element such as suture, suture wire, wire or equivalent (not shown) can extend from a separation point at or near the penetrating element and terminate at a needle (not shown). Staple devices according to embodiments of the invention preferably are made of a biocompatible material, e.g. titanium, stainless steel, 20 nitinol, etc. Such staple devices preferably are malleable, for crimping into a closed position with a closing or crimping device, and/or are constructed of a memory material or other material that can be induced to deform into a desired configuration, e.g. a closed configuration. In the case of a memory material such as nitinol, the staple device can be formed in the closed position, sprung open 25 and delivered to the tissues, and then released from an e.g. delivery system after

springing back into a closed position. Such memory material eliminates the need for complicated crimping tools, which are complex and potentially damaging to delicate tissues during deployment due to their bulk and the ergonomics required to deploy them.

5 Spacing elements according to embodiments of the invention substantially prevent the apposing evertng platforms from crushing the cut ends of the vascular or other structures in the anastomosis, by keeping the evertng platforms slightly separated from one another in the closed position of the staple. Thus, any tissue in the anastomosis is approximated without being strangulated, 10 consistent with the surgical adage: "approximate, don't strangulate." If the eversion platforms were not kept slightly apart in the closed position, the tissue engaged between them could be crushed to death (necrose) closer to the lumen. With particular reference to e.g. Figures 5-7, but also with reference to other embodiments, tissue is engaged in the closed staple with more tissue 15 compression at the spacing elements and less tissue compression at the offset evertng platforms. Tissue engagement and compression in this manner accomplishes approximation without strangulation, ensuring viability of the tissue at the most critical point of the anastomosis, namely, the juncture of the two structures being anastomosed.

20 As referenced above, a plurality of the evertng staple devices described herein can be used in a single anastomosis, or a single device can be used. In the case of a single device, one or more sutures, one or more conventional or other type of staples, one or more areas of biological adhesive and/or other means additionally can be used, e.g. at least directly opposite one or each staple device 25 described herein, to complete the anastomosis. In the case of multiple staple

devices, whatever the type, it is desirable to space each device at selected points around the circumference of the anastomosis, e.g. at quadrant points (e.g. 12 o'clock, 3 o'clock, 6 o'clock and 9 o'clock positions) or half points (e.g. 12 o'clock and 6 o'clock positions).

5 Non-everting staple devices can be used at e.g. the heel and toe of an anastomosis formed using a longitudinal (i.e. in the direction of blood flow) opening in the artery or other recipient vessel, and everting staple devices can be used along the sides of the anastomosis between the heel and toe. Such use of non-everting staple devices would reduce the possibility of compromising the 10 circumference of the recipient vessel at the heel and toe. Use of adhesive, suture, or other connection devices or methods in addition to or instead of everting staples can also reduce possible complications and/or the number of everting staple devices needed to form an anastomosis. Staple devices all of the 15 everting type can be used more readily with a transverse (i.e. in the direction perpendicular to blood flow) opening in the recipient vessel.

Of course, each staple can be appropriately dimensioned to suit a particular patient, surgical procedure, surgical environment or other factor. Devices of different sizes can be used in the same anastomosis, if desired.

In use, according to one embodiment, the surgeon or other medical 20 professional penetrates the walls of the vascular or other anatomical structures using one or more needles attached to e.g. suture or other material. The suture, in turn, is attached at or near e.g. a penetrating element of the staple device or to another suitable point at, on, or in association with the staple device. According 25 to alternative embodiments, the staple is a free-standing staple and/or includes delivery mechanisms other than suture and one or more needles.

While the invention has been described with respect to specific embodiments, the invention should not be considered limited to the specific embodiments illustrated and described herein. For example, embodiments of the invention apply not only to anastomoses of vascular structures in e.g. minimally invasive thoracic surgical situations, but also to conventional surgical techniques and to anatomical structures other than vascular structures. Embodiments of the invention apply to anastomosing prosthetic tubular grafts to vascular structures or to each other. Specific features described with respect to one embodiment are also to be considered useable with the other embodiments disclosed and contemplated herein. Other modifications and changes are readily discernible from the specification and will be apparent to those of ordinary skill.

WHAT IS CLAIMED IS:

1. A surgical staple for use in creating an everted anastomosis of at least two anatomical structures, the staple comprising a staple body constructed for bending and at least two evertting elements connected to the staple body, the evertting elements protruding from the staple body, the staple body and the evertting elements being constructed and arranged so that when the staple body is bent around cut ends of the anatomical structures, the evertting elements are in an apposed relationship and the inner layers of the anatomical structures are held together between the evertting elements to form the everted anastomosis.
2. The staple of claim 1, further comprising at least two spacing elements connected to the staple body, the spacing elements being constructed and arranged so that when the staple body is bent around the cut ends of the anatomical structures, the spacing elements are in an apposed relationship and the cut ends of the anatomical structures are held together between the spacing elements.
3. The staple of claim 2, wherein the distance between the apposed spacing elements is less than the distance between the apposed evertting elements.
4. The staple of claim 2, wherein the apposed evertting elements are closer than the apposed spacing elements to the lumen of the anastomosed anatomical structures.

5. The staple of claim 2, wherein the apposed spacing elements define an inner radius of the bent staple and the apposed evertting elements define an outer radius of the bent staple, the radii being taken from the bend in the staple.
6. The staple of claim 2, wherein the spacing elements comprise collars disposed around the staple body.
7. The staple of claim 6, wherein at least one of the collars is constructed to slide along the staple body into apposed relationship with the other of the collars.
8. The staple of claim 2, wherein one of the spacing elements is disposed along the staple body and the other of the spacing elements is disposed adjacent the staple body.
9. The staple of claim 2, wherein the spacing elements are disposed closer to the middle of the staple than the evertting elements.
10. The staple of claim 1, wherein the evertting elements are disposed at the very ends of the staple body.
11. The staple of claim 1, further comprising a penetrating element constructed to penetrate the ends of the anatomical structures when the evertting elements are in the apposed relationship.

12. The staple of claim 11, wherein the penetrating element is disposed at one end of the staple body.
13. The staple of claim 1, wherein the evertting elements are offset from each other transversely with respect to the staple body.
14. The staple of claim 13, wherein the end of one of the evertting elements is aligned with the staple body; further wherein one other of the evertting elements is centered on the staple body.
15. The staple of claim 13, wherein one of the evertting elements moves laterally with respect to the staple body as the staple is bent, such that the direction of closure of the staple is at an angle to the staple body.
16. The staple of claim 1, wherein the staple body is outside the lumen of the anastomosed anatomical structures when the evertting elements are apposed.
17. The staple of claim 1, constructed such that the lumen of the anastomosed anatomical structures is free of exposure to the staple and to cut tissue edges of the anatomical structures.
18. The staple of claim 1, wherein the staple is formed of a memory metal.
19. A surgical staple for use in creation of an everted anastomosis between at least two anatomical structures, the staple comprising means for bending and

means for evertng connected to and protruding from the means for bending, the means for bending and the means for evertng being constructed and arranged so that when the means for bending is bent around cut ends of the anatomical structures, the means for evertng forms an apposed relationship and the inner layers of the anatomical structures are held together between the means for evertng to form the everted anastomosis.

20. The staple of claim 19, further comprising means for spacing connected to the staple body, the means for spacing being constructed and arranged so that when the staple body is bent around the cut ends of the anatomical structures, the means for spacing are in an apposed relationship and the cut ends of the anatomical structures are held together between the spacing elements.

21. The staple of claim 19, wherein the means for evertng and the means for spacing are disposed on opposite ends of the staple.

22. A surgical staple constructed to join at least two anatomical structures, the staple comprising:

 a first evertng platform disposed at a first end of the staple, the first evertng platform being centered at the first end of the staple;

 a second evertng platform disposed at a second end of the staple, the second evertng platform being connected at one end thereof to the second end of the staple, the first and second evertng platforms being constructed to form an everted anastomosis of the anatomical structures;

a first spacing element disposed at the first end of the staple, the first spacing element being disposed adjacent the longitudinal axis of the staple; a second spacing element disposed at the second end of the staple, the second spacing element being disposed along the longitudinal axis of the staple; and

a penetrating element for penetrating the anatomical structures, the penetrating element being disposed at the first end of the staple along the longitudinal axis of the staple.

23. The staple of claim 22, wherein the staple is constructed to bend to a closed configuration in which the second evertng platform is disposed parallel to the first evertng platform and in which the first and second evertng platforms are disposed on the same side of the penetrating element.

24. The staple of claim 23, wherein in the closed configuration the first and second spacing elements are apposed.

25. The staple of claim 24, wherein in the closed configuration the first and second spacing elements and the penetrating element define an inner radius of the staple; further wherein in the closed configuration the first and second evertng platforms define an outer radius of the staple.

26. The staple of claim 23, wherein in the closed configuration the staple is bent into a "U" shape to form two legs and a bend, the legs of the staple being offset from each other in two dimensions.

27. A method of closing a surgical staple, the method comprising:

moving a first evertting platform disposed at one end of the staple to be parallel to and adjacent to a second evertting platform disposed at an opposite end of the staple;

moving a first spacing element disposed at said one end of the staple to be apposed to a second spacing element disposed at said opposite end of the staple; and

disposing the first and second spacing elements adjacent to and generally parallel to a penetrating element of the staple, the penetrating element being constructed to penetrate anatomical structure.

28. The method of claim 27, further comprising moving the first evertting platform in a direction of closure to its disposition parallel to and adjacent to the second evertting platform, the direction of closure being disposed at an angle to a longitudinal axis of the staple extending between the first evertting platform and the second evertting platform.

29. The method of claim 27, wherein the first and second evertting platforms are moved from an offset configuration, in which the first and second evertting platforms are offset from each other along their longitudinal axes, to a non-offset configuration, in which the first and second evertting platforms are aligned with each other.

30. A device for use in creating an anastomosis of two anatomical structures, the anastomosis defining a central lumen, the device comprising:

penetration structure for penetrating and holding both anatomical structures at the anastomosis; and

eversion structure for everting an end of at least one of the anatomical structures, the eversion structure being disposed closer to the lumen of the anastomosis than the penetration structure, the eversion structure being constructed to accommodate the anatomical structures.

31. The device of claim 30, wherein the device is constructed from a memory metal such that a crimping tool is unnecessary to create the anastomosis.

32. The device of claim 30, further comprising spacing structure for providing space within the eversion structure to minimize tissue necrosis in the anatomical structures.

33. The device of claim 30, wherein the device is a surgical staple, further wherein the penetration structure defines an inner radius of the staple and the eversion structure defines an outer radius of the staple, placement of the inner radius and outer radius causing the staple to be excluded from the lumen of the anastomosis.

34. The device of claim 30, wherein the device is a surgical staple having a staple body, further wherein the eversion structure comprises two everting platforms protruding from the staple body.

35. The device of claim 30, wherein the device is an evertting surgical staple, the evertting surgical staple being in combination with at least one non-evertting surgical staple to form the anastomosis.

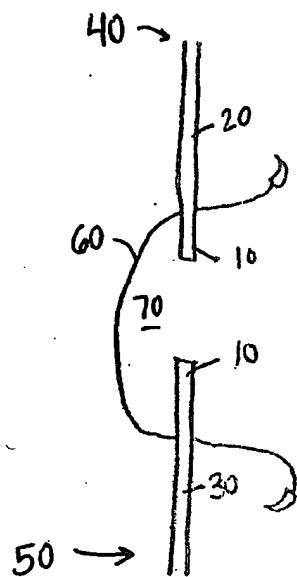


Fig. 1A

PRIOR ART

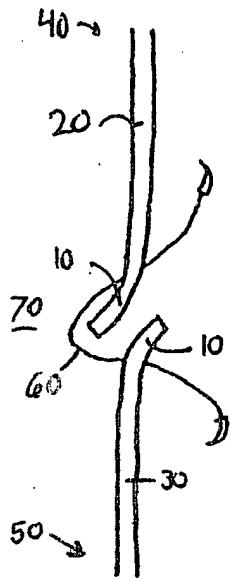


Fig. 2A

PRIOR ART

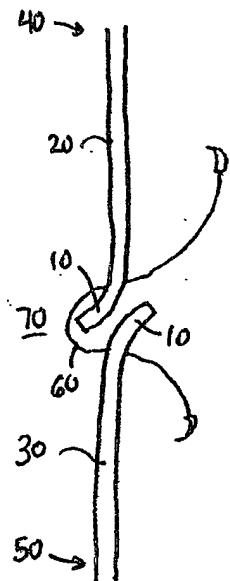


Fig. 3A

PRIOR ART

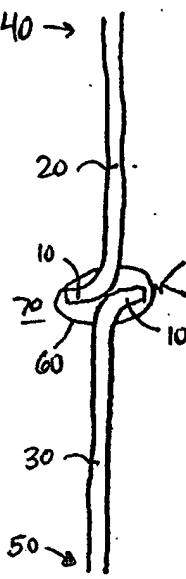


Fig. 4A

PRIOR ART

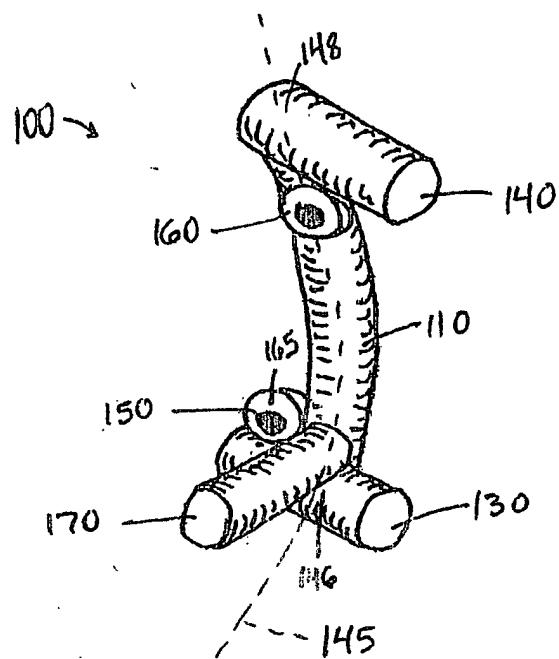


Fig. 2

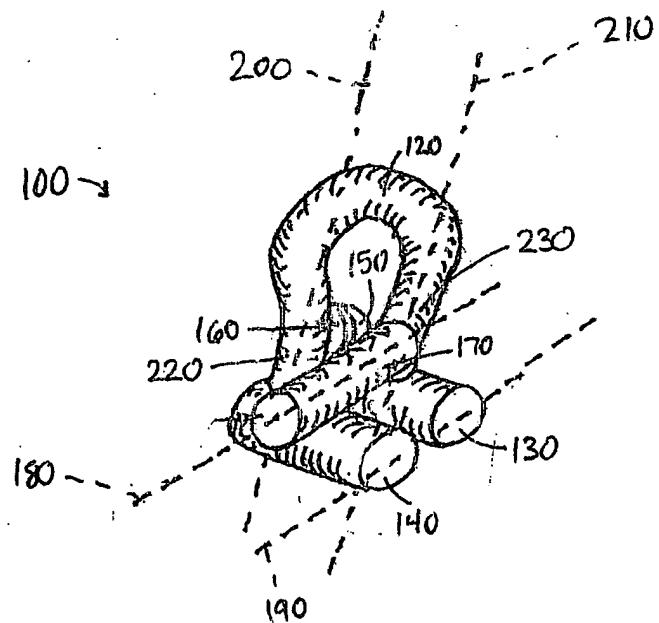


Fig. 3

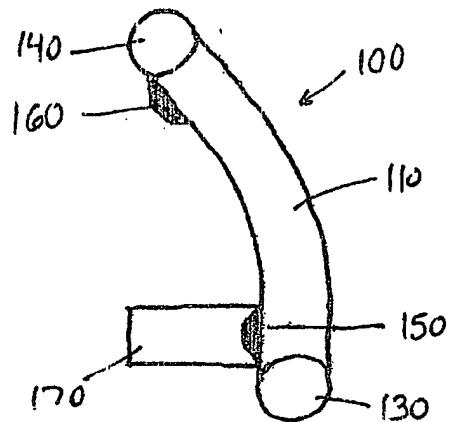


Fig. 4

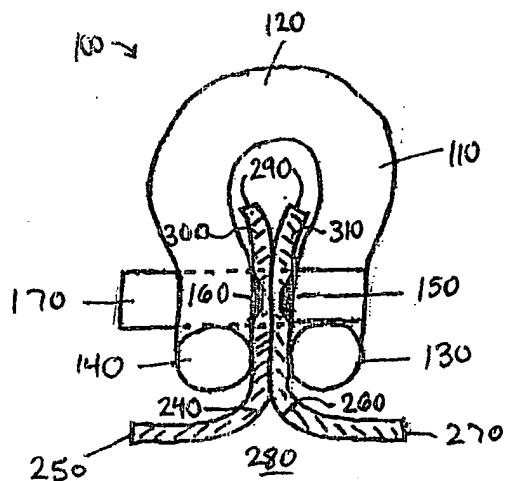


Fig. 5

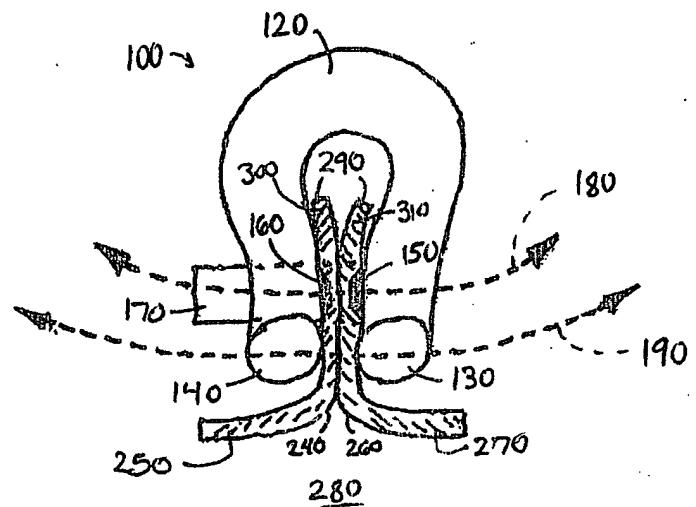


Fig. 6

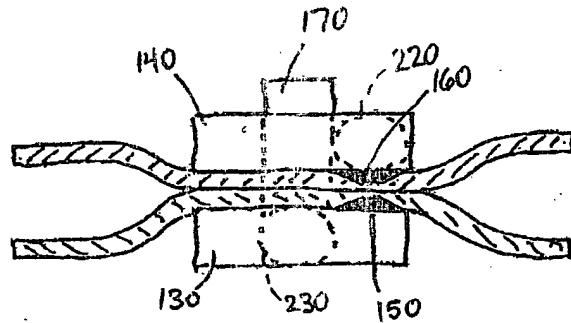


Fig. 7

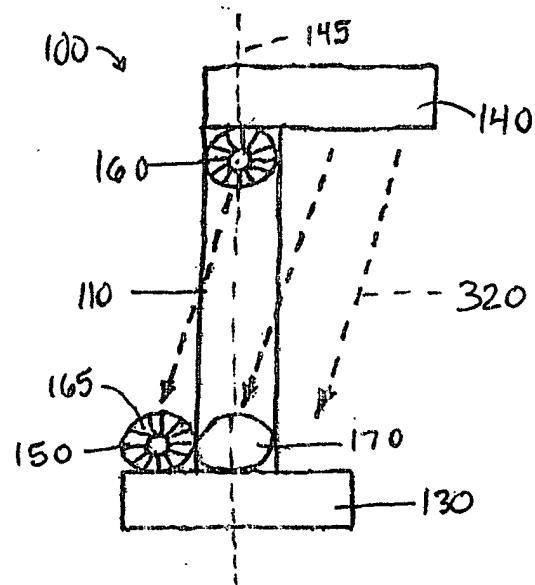


Fig. 8

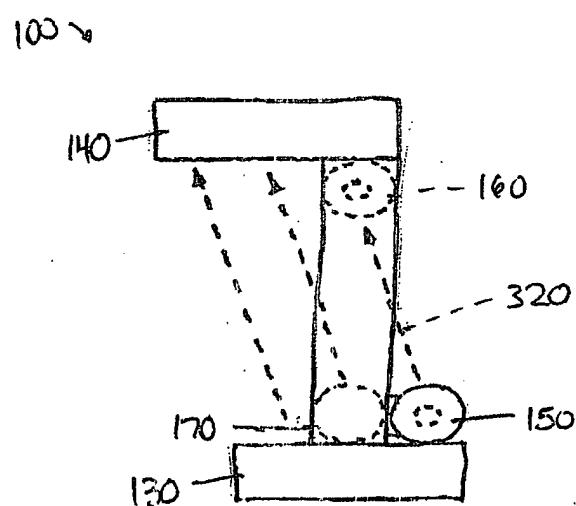


Fig. 9

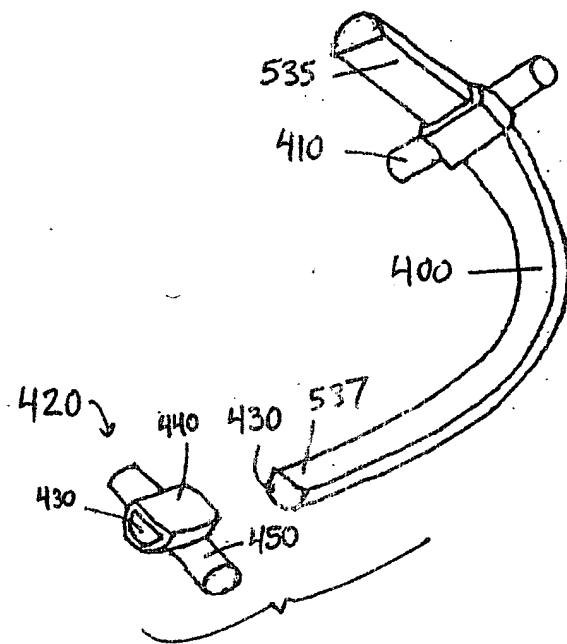


Fig. 10

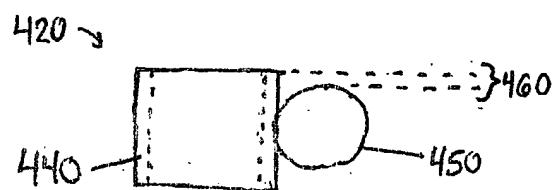


Fig. 11

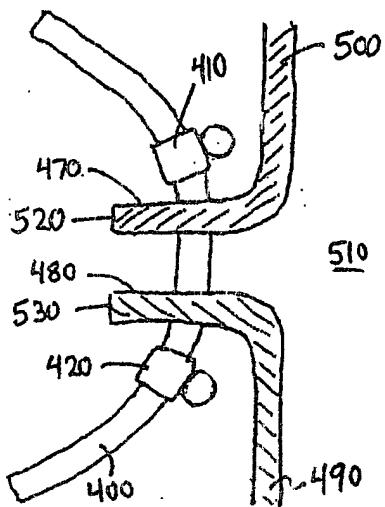


Fig. 12

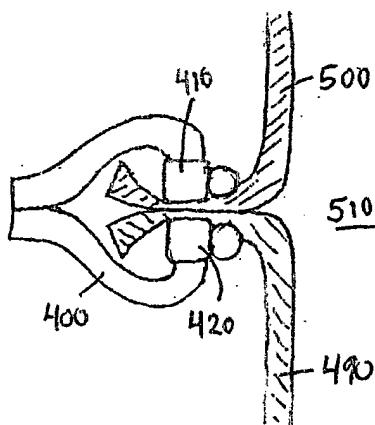


Fig. 13

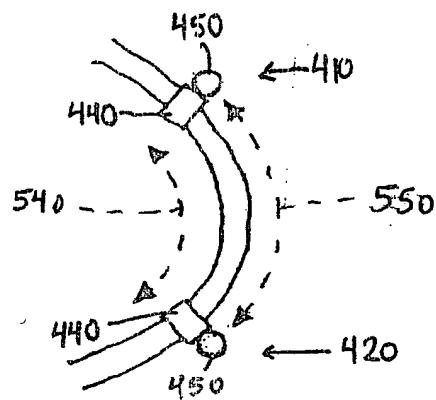


Fig. 14

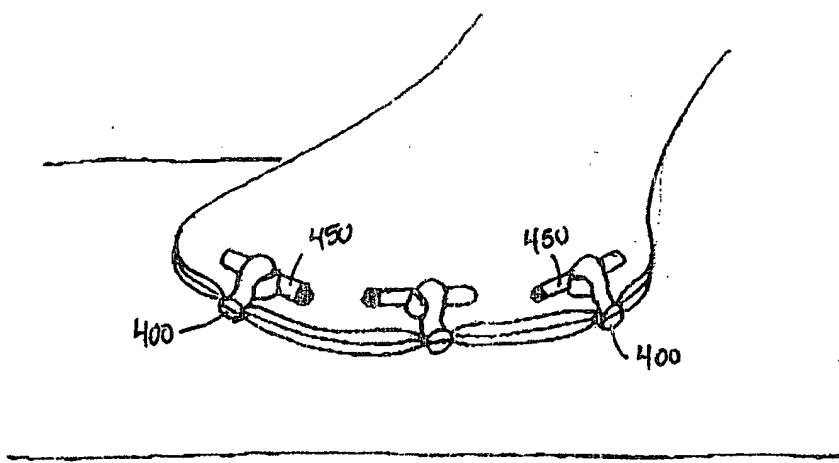


Fig. 15

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 01/07337A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61B17/064

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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A	WO 98 30153 A (ELECTIVE VASCULAR INTERVENTION) 16 July 1998 (1998-07-16) abstract; figures 1,5	22
X	WO 98 30152 A (ELECTIVE VASCULAR INTERVENTION) 16 July 1998 (1998-07-16) abstract; figures 1,2	1,10,19, 30
A	WO 99 62406 A (COALESCENT SURGICAL INC ;GANDIONCO ISIDRO MATIAS (US); HO LIEM (US) 9 December 1999 (1999-12-09) abstract; figure 2	1-26, 30-35
		1-26, 30-35
		-/-

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

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Date of the actual completion of the international search

27 September 2001

Date of mailing of the International search report

05/10/2001

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Hansen, S

INTERNATIONAL SEARCH REPORT

Int'l Application No.
PCT/US 01/07337

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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